It is hereby certified that this photocopy is a true copy of the original document shown to me.
Uppsala, Sweden, March 1, 2018
Ex officio:

Olof Wijk
Notary Public

NOTARIUS PUBLICUS I UPPSALA

The Ministry for Foreign Affairs in Stockholm,
hereby certifies that
Mr. Olof Wijk
Notary Public of Uppsala
has issued and signed the foregoing
attestation in his official capacity.

Stockholm 5 March 2018

Nr. 4856

Fee Paid 180 SEK

Essie Persson

CHỨNG NHẬN/HỢP PHÁP HÓA LÃNH SỰ
CONSULAR AUTHENTICATION

1. Quốc gia *Country* Việt Nam / Vietnam
Giấy tờ, tài liệu này *This public document*
2. do Ông (Bà) Essie Persson ký *has been signed by*
3. với chức danh *acting in the capacity of*
4. và con dấu của Bộ Ngoại Giao Thụy Điển *bears the seal/stamp of* Swedish Ministry for Foreign Affairs

được chứng nhận/hợp pháp hóa lãnh sự *Certified*

5. tại *at* Stockholm 6. ngày *the* 05/02/2018
7. Cơ quan cấp *by* DSQ VN / Vn Embassy
8. Số: 111/2018 *Nº*

TL. ĐẠI SỨ / FOR THE AMBASSADOR
Bí thư thứ Hai / Second Secretary

Phạm Thị Quyên

kibion

EG-KONFORMITÄTSERKLÄRUNG
EC DECLARATION OF CONFORMITY
gemäß Anhang III der Richtlinie 98/79/EC über In-vitro-Diagnostika
according to Annex III of the IVD Medical Device Directive 98/79/EC

Hersteller / Manufacturer:	Kibion GmbH
Branche / Business sector:	Atemtests / Breath tests
Anschrift / Address:	Haferwende 31 D-28357 Bremen, Germany
IVD-Medizinprodukt / IVD-Medical Device:	Kibion Dynamic Pro
Klassifizierung gemäß 98/79/EG / Classification according to 98/79/EC:	Sonstiges Produkt (nicht im Anhang II, nicht zur Eigenanwendung) Other product (not in Annex II, not for self-testing)
Nomenklatursystem & -Code/ Code and Term:	EDMS 26-09. Kategorie 06

Angaben zum Produkt / Device identification		
Produkt / Product	Artikel Nr./ REF no.	Beschreibung, Version etc. Description, version, etc.
Kibion® Dynamic Pro	8032	Erweiterungseinheit für das Kibion® Dynamic Base (Infrarot Isotopen Analysator für 13C-Atemgas) mit 16 Anschlüssen Port extension unit for IRIS Dynamic base (Infrared Isotop Analyser for 13C breath tests) with 16 ports

Wir erklären in alleiniger Verantwortung, dass die Medizinprodukte allen anwendbaren Anforderungen der Richtlinie 98/79/EC für in-vitro-Diagnostika bzw. des Gesetzes über Medizinprodukte (MPG) entsprechen.

Das Produkt ist CE-zertifiziert durch Kibion GmbH.

Nicht von uns ermächtigte Produkt-änderungen machen diese Erklärung ungültig.

We declare, in sole responsibility, that the concerned medical devices comply with the applicable requirements of the IVD-directive 98/79/EC and with the German law concerning Medical Devices (MPG) respectively.

The product is CE-marked by Kibion GmbH.

Any modification of the devices, not authorised by us, will invalidate this declaration.

Ort und Datum / Place and Date	UPPSALA 2017-03-10
Unterschrift / Signature	
Name / Printed name	Petter Bäckgren
Funktion / Position	Geschäftsführer / CEO

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Rev04, Br-219-03, March 2017, Kibion AB